## Claims

1. A compound of the formul

 $\mathbb{R}^{1} = \mathbb{R}^{1} \times \mathbb{R}^{2} \times \mathbb{R}^{2}$ 

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wherein R is lower-alkyl, R<sup>1</sup> is halogen, R<sup>2</sup> is  $C_1-C_{12}$ -alkyl, R<sup>3</sup> is hydroxy, lower-alkoxy, lower-alkylcarbonyloxy, lower-alkylcarbonyloxy, lower-alkylaminocarbonyloxy, arylaminocarbonyloxy or aryl-lower-alkylaminocarbonyloxy, X is  $C_1-C_{18}$ -alkylene which optionally can be interrupted by 1.4-phenylene or interrupted or lengthened by 1.4-cyclohexylene, A is di- or tri-substituted 2-imidazolyl attached via an ethylene group or a substituted or unsubstituted heterocycle selected from the group consisting of benzimidazolyl, benzimidazolonyl, imidazo[4,5-c]pyridinyl, imidazo-[4,5-c]pyridinonyl, benzthiazolyl, benzodiazepine-2,5-dion-1-yl and pyrrolo[2,1-c][1,4]benzodiazepine-5,11-dion-10-yl and n is the number O or 1,

in the form of a racemate or an optical antipode, an N-oxide, or a pharmaceutically usable acid addition salt thereof.

2. A compound in accordance with claim 1, wherein R is isopropyl.

3. A compound in accordance with claim 2, wherein R<sup>3</sup> is hydroxy, lower-alkylcarbonyloxy, lower-alkoxy-lower-alkylcarbonyloxy or lower-alkylaminocarbonyloxy.

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4. A compound in accordance with claim 3, wherein  ${\tt R}^3$  is isobutyryloxy, methoxyacetyloxy or butylaminocarbonyloxy.

5. A compound in accordance with claim 1, wherein r is the number 1.

6. A compound in accordance with claim 1, wherein R<sup>1</sup> is fluorine.

7. A compound in accordance with claim 1, wherein R<sup>2</sup> is methyl.

8. A compound in accordance with claim 1. wherein x 15 is C<sub>3</sub>-C<sub>7</sub>-alkylene.

9. A compound in accordance with claim 8, wherein X is propylene, butylene, pentamethylene or hexamethylene.

10. A compound in accordance with claim 1, wherein A is 2-benzimidazolyl, 2-benzthiazolyl, 1-methyl-2-benzimidazolyl, 1-dodecyl-2-benzimidazolyl, benzimidazolonyl, 2,3,4,5-tetrahydro-4-methylbenzodiazepine-2,5-dion-1-yl, 6-chloro-2,3,11,11a-tetrahydro-pyrrolo[2,1-c][1,4]benzo-diazepine-5,11-dion-10-yl or 1-methyl-4,5-diphenyl-2-imidazolyl.

11. A compound in accordance with claim 10, wherein A
is 2-benzimidazolyl or 2-benzthiazolyl.

12. A compound in accordance with claim 2, wherein R is isopropyl, R is hydroxy, isobutyryloxy, methoxy-acetyloxy or butylaminocarbonyloxy, R is fluorine, R is methyl, X is propylene, butylene, pentamethylene or hexamethylene, A is 2-benzimidazolyl or 2-benzthiazolyl and n is the number 1.

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13. A compound in accordance with claim 2.)

2-[2-[[3-(2-benzimidazolyl)propyl]methylamino]
ethyl]-6-fluoro-1.2.3.4-tetrahydro-lα-isopropyl-2α
-naphthyl methoxyacetate.

14. A compound in accordance with claim 1.

[1S.2S]-2-[2-[[5-(2-benzthiazolyl)pentyl]methylamino]ethyl]-6-fluoro-1,2,3,4-tetrahydro-1-isopropyl-2-naphthyl methoxyacetate.

15. A compound in accordance with claim [15.25]-2-[2-[[3-(2-benzimidazolyl)propyl]methylamino]ethyl]-6-fluoro-1,2,3,4-tetrahydro-1-isopropyl-2-naphthyl methoxyacetate.

16. A composition with calcium antagonistic activity comprising an effective amount of a compound of the formula

R R 3
N-(X) n-A

wherein R is lower-alkyl, R<sup>1</sup> is halogen, R<sup>2</sup> is  $C_1-C_{12}$ -alkyl, R<sup>3</sup> is hydroxy, lower-alkoxy, lower-alkylcarbonyloxy, lower-alkylcarbonyloxy, lower-alkylaminocarbonyloxy, arylaminocarbonyloxy or aryl-lower-alkylaminocarbonyloxy, X is  $C_1-C_{18}$ -alkylene which optionally can be interrupted by 1.4-phenylene or interrupted or lengthened by 1.4-cyclohexylene, A is di- or tri-substituted 2-imidazolyl attached via an ethylene group or a substituted or unsubstituted heterocycle selected from the group consisting of benzimidazolyl, benzimidazolonyl, imidazo[4.5-c]pyridinyl, imidazo-[4.5-c]pyridinonyl, benzodiazepine-2.5-

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-dion-1-yl and pyrrolo[2.1-c][1.4]benzodiazepine-5.11-dion-10-yl and n is the number o or 1. in the form of a racemate or an optical antipode, an N-oxide, or a pharmaceutically usable acid addition salt thereof, and a pharmaceutically inert excipient.

- 17. A composition in accordance with claim 16.

  wherein R is isopropyl, R is hydroxy, isobutyryloxy,
  methoxyacetyloxy or butylaminocarbonyloxy, R is

  10 fluorine, R is methyl, X is propylene, butylene, pentamethylene or hexamethylene, A is 2-benzimidazolyl or
  2-benzthiazolyl and n is the number 1.
- 18. A composition in accordance with claim 17,

  wherein the compound of formula I is [15,25]-2-[2-[[3--(2-benzimidazolyl)propyl]methylamino]ethyl]-6-fluoro--1,2,3,4-tetrahydro-1-isopropyl-2-naphthyl methyoxy-acetate or its racemate.
  - 19. A method of treating or preventing angina pectoris, ischaemia, arrhythmias, high blood pressure and cardiac insufficiency which comprises administering to a warm-blooded animal in need of such treatment, an effective amount of a compound of the formula \(\cap\$

<sub>R</sub>1/

 $I = (X)_{n} - I$ 

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wherein R is lower-alkyl, R<sup>1</sup> is halogen, R<sup>2</sup> is  $C_1^{-C_{12}^{-}}$  alkyl, R<sup>3</sup> is hydroxy, lower-alkoxy, lower-alkylcarbonyloxy, lower-alkylcarbonyloxy, lower-alkylcarbonyloxy, lower-alkylaminocarbonyloxy, arylaminocarbonyloxy or aryl-lower-alkylaminocarbonyloxy, X is  $C_1^{-C_{18}^{-}}$  alkylene which optionally can be interrupted by

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1.4-phenylene or interrupted or lengthened by
1.4-cyclohexylene. A is di- or tri-substituted
2-imidazolyl attached via an ethylene group or a
substituted or unsubstituted heterocycle selected from
the group consisting of benzimidazolyl, benzimidazolonyl, imidazo[4.5-c]pyridinyl, imidazo[4.5-c]pyridinonyl, benzthiazolyl, benzodiazepine-2.5-dion-1-yl and pyrrolo[2.1-c][1.4]benzodiazepine-5.11-dion-10-yl and n is the number 0 or 1,

in the form of a racemate or an optical antipode, an N-oxide, or a pharmaceutically usable acid addition salt thereof.

20. A method in accordance with claim 19, wherein R is isopropyl, R is hydroxy, isobutyryloxy, methoxy-acetyloxy or butylaminocarbonyloxy, R is fluorine, R is methyl, X is propylene, butylene, pentamethylene or hexamethylene, A is 2-benzimidazolyl or 2-benzthiazolyl and n is the number 1.

21. A method in accordance with claim 20, wherein the compound of formula I is [18,28]-2-[2-[[3-(2-benzimida-zolyl)propyl]methylamino]ethyl]-6-fluoro-1,2,3,4-tetra-hydro-1-isopropyl-2-naphthyl methoxyacetate or its racemate.

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